

# Jantina De Vries

## List of Publications by Year in descending order

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Version: 2024-02-01

104  
papers

3,678  
citations

172457

29  
h-index

155660

55  
g-index

109  
all docs

109  
docs citations

109  
times ranked

4055  
citing authors

#	ARTICLE	IF	CITATIONS
1	Should Feedback of Individual Results be Integrated into the Consent Process in African Genomics? Participants' Views from an HIV-TB Genomics Research Project in Botswana. <i>AJOB Empirical Bioethics</i> , 2022, 13, 48-56.	1.6	2
2	COVID-19 vaccine trials with children: ethics pointers. <i>BMJ Global Health</i> , 2022, 7, e007466.	4.7	5
3	Guideline for feedback of individual genetic research findings for genomics research in Africa. <i>BMJ Global Health</i> , 2022, 7, e007184.	4.7	8
4	Towards equitable genomics governance in Africa: Guiding principles from theories of global health governance and the African moral theory of Ubuntu. <i>Bioethics</i> , 2022, 36, 411-422.	1.4	3
5	Considerations of Autonomy in Guiding Decisions around the Feedback of Individual Genetic Research Results from Genomics Research: Expectations of and Preferences from Researchers in Botswana. <i>Global Health, Epidemiology and Genomics</i> , 2022, 2022, 1-7.	0.8	5
6	Social and Epistemic Justice: Are We Really Including Africa in the Bioethics Discourse?. <i>American Journal of Bioethics</i> , 2022, 22, 30-32.	0.9	0
7	Unpacking race and ethnicity in African genomics research. <i>Nature Reviews Genetics</i> , 2022, 23, 455-456.	16.3	1
8	Tailored community engagement to address the genetics diversity gap. <i>Med</i> , 2022, 3, 369-370.	4.4	1
9	Mapping experiences and perspectives of equity in international health collaborations: a scoping review. <i>International Journal for Equity in Health</i> , 2021, 20, 28.	3.5	22
10	Negotiating political power and stigma around fragile X Syndrome in a rural village in Cameroon: A tale of a royal family and a community. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2021, 9, e1615.	1.2	0
11	Genomics governance: advancing justice, fairness and equity through the lens of the African communitarian ethic of Ubuntu. <i>Medicine, Health Care and Philosophy</i> , 2021, 24, 377-388.	1.8	12
12	The role of causal knowledge in stigma considerations in African genomics research: Views of South African Xhosa people. <i>Social Science and Medicine</i> , 2021, 277, 113902.	3.8	2
13	POPIA Code of Conduct for Research (with corrigendum). <i>South African Journal of Science</i> , 2021, 117, .	0.7	8
14	Explanatory models for the cause of Fragile X Syndrome in rural Cameroon. <i>Journal of Genetic Counseling</i> , 2021, 30, 1727-1736.	1.6	3
15	Evaluating Community Engagement Strategies to Manage Stigma in Two African Genomics Studies Involving People Living with Schizophrenia or Rheumatic Heart Disease. <i>Global Health, Epidemiology and Genomics</i> , 2021, 2021, 1-9.	0.8	3
16	A scoping review of considerations and practices for benefit sharing in biobanking. <i>BMC Medical Ethics</i> , 2021, 22, 102.	2.4	10
17	Systematic Review of the Economic Evaluation of Returning Incidental Findings in Genomic Research. <i>Frontiers in Public Health</i> , 2021, 9, 697381.	2.7	4
18	Genome-wide association studies. <i>Nature Reviews Methods Primers</i> , 2021, 1, .	21.2	529

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19	Association of Novel Locus With Rheumatic Heart Disease in Black African Individuals. <i>JAMA Cardiology</i> , 2021, 6, 1000.	6.1	18
20	Toward better governance of human genomic data. <i>Nature Genetics</i> , 2021, 53, 2-8.	21.4	31
21	Considering equity in global health collaborations: A qualitative study on experiences of equity. <i>PLoS ONE</i> , 2021, 16, e0258286.	2.5	14
22	Participants' Preferences and Reasons for Wanting Feedback of Individual Genetic Research Results From an HIV-TB Genomic Study: A Case Study From Botswana. <i>Journal of Empirical Research on Human Research Ethics</i> , 2021, 16, 525-536.	1.3	6
23	Engaged genomic science produces better and fairer outcomes: an engagement framework for engaging and involving participants, patients and publics in genomics research and healthcare implementation. <i>Wellcome Open Research</i> , 2021, 6, 311.	1.8	6
24	Use of broad consent and related procedures in genomics research: Perspectives from research participants in the Genetics of Rheumatic Heart Disease (RHDGen) study in a University Teaching Hospital in Zambia. <i>Global Bioethics</i> , 2020, 31, 184-199.	1.5	17
25	What constitutes good ethical practice in genomic research in Africa? Perspectives of participants in a genomic research study in Uganda. <i>Global Bioethics</i> , 2020, 31, 169-183.	1.5	20
26	Returning incidental findings in African genomics research. <i>Nature Genetics</i> , 2020, 52, 17-20.	21.4	36
27	Do solidarity and reciprocity obligations compel African researchers to feedback individual genetic results in genomics research?. <i>BMC Medical Ethics</i> , 2020, 21, 112.	2.4	24
28	Global citizen deliberation on genome editing. <i>Science</i> , 2020, 369, 1435-1437.	12.6	47
29	Benefit Sharing for Human Genomics Research: Awareness and Expectations of Genomics Researchers in Sub-Saharan Africa. <i>Ethics &amp; Human Research</i> , 2020, 42, 14-20.	0.9	15
30	Exploring how a genetic attribution to disease relates to stigma experiences of Xhosa patients with schizophrenia in South Africa. <i>Social Psychiatry and Psychiatric Epidemiology</i> , 2020, 55, 1679-1686.	3.1	8
31	Lived Experiences of Fragile X Syndrome Caregivers: A Scoping Review of Qualitative Studies. <i>Frontiers in Neurology</i> , 2020, 11, 128.	2.4	4
32	How Should Biobanking Be Governed in Low-Resource Settings?. <i>AMA Journal of Ethics</i> , 2020, 22, E156-163.	0.7	8
33	Stigma in African genomics research: Gendered blame, polygamy, ancestry and disease causal beliefs impact on the risk of harm. <i>Social Science and Medicine</i> , 2020, 258, 113091.	3.8	15
34	Informed consent in genomic research and biobanking: taking feedback of findings seriously. <i>Global Bioethics</i> , 2020, 31, 200-215.	1.5	18
35	The ethics of neurogenetics research in Africa: Considerations and guidelines. , 2020, , 81-94.		0
36	The governance of genomic biobank research in Africa: reframing the regulatory tilt. <i>Journal of Law and the Biosciences</i> , 2020, 7, 1-18.	1.6	13

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37	Cascade Testing for Fragile X Syndrome in a Rural Setting in Cameroon (Sub-Saharan Africa). <i>Genes</i> , 2020, 11, 136.	2.4	7
38	Providing a policy framework for responsible gene drive research: an analysis of the existing governance landscape and priority areas for further research. <i>Wellcome Open Research</i> , 2020, 5, 173.	1.8	12
39	Ethical considerations in genomic research in South Africa. <i>South African Medical Journal</i> , 2019, 109, 375.	0.6	6
40	Dissecting in silico Mutation Prediction of Variants in African Genomes: Challenges and Perspectives. <i>Frontiers in Genetics</i> , 2019, 10, 601.	2.3	25
41	T42EXPLORING WHETHER A GENETIC ATTRIBUTION TO DISEASE RELATES TO INTERNALISED STIGMA EXPERIENCES OF XHOSA PATIENTS WITH SCHIZOPHRENIA IN SOUTH AFRICA. <i>European Neuropsychopharmacology</i> , 2019, 29, S238-S239.	0.7	0
42	Engaging research ethics committees to develop an ethics and governance framework for best practices in genomic research and biobanking in Africa: the H3Africa model. <i>BMC Medical Ethics</i> , 2019, 20, 69.	2.4	23
43	Safeguarding the future of genomic research in South Africa: Broad consent and the Protection of Personal Information Act No. 4 of 2013. <i>South African Medical Journal</i> , 2019, 109, 468.	0.6	23
44	Consent insufficient for data release. <i>Science</i> , 2019, 364, 445-446.	12.6	9
45	Potential use of clinical polygenic risk scores in psychiatry – ethical implications and communicating high polygenic risk. <i>Philosophy, Ethics, and Humanities in Medicine</i> , 2019, 14, 4.	1.5	73
46	An optimistic vision for biosciences in South Africa: Reply to Thaldar et al. (2019). <i>South African Journal of Science</i> , 2019, 115, .	0.7	0
47	Does genetics matter for disease-related stigma? The impact of genetic attribution on stigma associated with rheumatic heart disease in the Western Cape, South Africa. <i>Social Science and Medicine</i> , 2019, 243, 112619.	3.8	8
48	Establishing a Multi-Country Sickle Cell Disease Registry in Africa: Ethical Considerations. <i>Frontiers in Genetics</i> , 2019, 10, 943.	2.3	14
49	The NeuroDev Study: Phenotypic and Genetic Characterization of Neurodevelopmental Disorders in Kenya and South Africa. <i>Neuron</i> , 2019, 101, 15-19.	8.1	10
50	Ethical and practical issues to consider in the governance of genomic and human research data and data sharing in South Africa: a meeting report. <i>AAS Open Research</i> , 2019, 2, 15.	1.5	5
51	Model framework for governance of genomic research and biobanking in Africa – a content description. <i>AAS Open Research</i> , 2018, 1, 13.	1.5	31
52	ASSAf consensus study on the ethical, legal and social implications of genetics and genomics in South Africa. <i>South African Journal of Science</i> , 2018, 114, .	0.7	6
53	Genomics research in Africa and its impact on global health: insights from African researchers. <i>Global Health, Epidemiology and Genomics</i> , 2018, 3, e12.	0.8	16
54	H3Africa: current perspectives. <i>Pharmacogenomics and Personalized Medicine</i> , 2018, Volume 11, 59-66.	0.7	101

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55	Predictors of consent to cell line creation and immortalisation in a South African schizophrenia genomics study. <i>BMC Medical Ethics</i> , 2018, 19, 72.	2.4	6
56	Community engagement in global health research that advances health equity. <i>Bioethics</i> , 2018, 32, 454-463.	1.4	41
57	Model framework for governance of genomic research and biobanking in Africa – a content description. <i>AAS Open Research</i> , 2018, 1, 13.	1.5	44
58	Regulation of genomic and biobanking research in Africa: a content analysis of ethics guidelines, policies and procedures from 22 African countries. <i>BMC Medical Ethics</i> , 2017, 18, 8.	2.4	76
59	Accessing Biospecimens from the H3Africa Consortium. <i>Biopreservation and Biobanking</i> , 2017, 15, 95-98.	1.0	15
60	Psychiatric Genomics: Ethical Implications for Public Health in Lower- and Middle-Income Countries. <i>American Journal of Bioethics</i> , 2017, 17, 17-19.	0.9	3
61	Stakeholder Perspectives on Public Health Genomics Applications for Sickle Cell Disease: A Methodology for a Human Heredity and Health in Africa (H3Africa) Qualitative Research Study. <i>OMICS A Journal of Integrative Biology</i> , 2017, 21, 323-332.	2.0	7
62	“I passed the test!” Evidence of diagnostic misconception in the recruitment of population controls for an H3Africa genomic study in Cape Town, South Africa. <i>BMC Medical Ethics</i> , 2017, 18, 12.	2.4	17
63	Developing the science and methods of community engagement for genomic research and biobanking in Africa. <i>Global Health, Epidemiology and Genomics</i> , 2017, 2, e13.	0.8	31
64	Ethical considerations in forensic genetics research on tissue samples collected post-mortem in Cape Town, South Africa. <i>BMC Medical Ethics</i> , 2017, 18, 66.	2.4	9
65	A Comprehensive Needs Assessment Tool for Planning RHD Control Programs in Limited Resource Settings. <i>Global Heart</i> , 2017, 12, 25.	2.3	13
66	Equity in international health research collaborations in Africa: Perceptions and expectations of African researchers. <i>PLoS ONE</i> , 2017, 12, e0186237.	2.5	67
67	Implications of direct-to-consumer whole-exome sequencing in South Africa. <i>South African Medical Journal</i> , 2016, 106, 139.	0.6	3
68	Evolving perspectives on broad consent for genomics research and biobanking in Africa. Report of the Second H3Africa Ethics Consultation Meeting, 11 May 2015. <i>Global Health, Epidemiology and Genomics</i> , 2016, 1, e13.	0.8	8
69	Obtaining informed consent for genomics research in Africa: analysis of H3Africa consent documents. <i>Journal of Medical Ethics</i> , 2016, 42, 132-137.	1.8	50
70	Ethical Challenges in Contemporary FASD Research and Practice. <i>Cambridge Quarterly of Healthcare Ethics</i> , 2016, 25, 726-732.	0.8	4
71	Deliberation to Promote Shared Sovereignty in Health Research: Four Questions to Clarify Goals, Methods, and Scope. <i>American Journal of Bioethics</i> , 2016, 16, 50-52.	0.9	3
72	Ethics review for international data-intensive research. <i>Science</i> , 2016, 351, 1399-1400.	12.6	44

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73	Broad Consent for Genomic Research and Biobanking: Perspectives from Low- and Middle-Income Countries. Annual Review of Genomics and Human Genetics, 2016, 17, 375-393.	6.2	59
74	Community engagement strategies for genomic studies in Africa: a review of the literature. BMC Medical Ethics, 2015, 16, 24.	2.4	87
75	The H3Africa policy framework: negotiating fairness in genomics. Trends in Genetics, 2015, 31, 117-119.	6.7	65
76	Addressing ethical issues in H3Africa research – the views of research ethics committee members. The HUGO Journal, 2015, 9, .	4.1	40
77	Exploring researchers' experiences of working with a researcher-driven, population-specific community advisory board in a South African schizophrenia genomics study. BMC Medical Ethics, 2015, 16, 45.	2.4	27
78	Would you terminate a pregnancy affected by sickle cell disease? Analysis of views of patients in Cameroon. Journal of Medical Ethics, 2014, 40, 615-620.	1.8	23
79	Enabling the genomic revolution in Africa. Science, 2014, 344, 1346-1348.	12.6	361
80	A perpetual source of DNA or something really different: ethical issues in the creation of cell lines for African genomics research. BMC Medical Ethics, 2014, 15, 60.	2.4	11
81	Knowing who to trust: exploring the role of "ethical metadata" in mediating risk of harm in collaborative genomics research in Africa. BMC Medical Ethics, 2014, 15, 62.	2.4	15
82	Ethical issues in genomic research on the African continent: experiences and challenges to ethics review committees. Human Genomics, 2014, 8, 15.	2.9	50
83	Consent and Community Engagement in Diverse Research Contexts: Reviewing and Developing Research and Practice. Journal of Empirical Research on Human Research Ethics, 2013, 8, 1-18.	1.3	123
84	Genomic sovereignty and the African promise: mining the African genome for the benefit of Africa. Journal of Medical Ethics, 2012, 38, 474-478.	1.8	68
85	ELSI 2.0 for Genomics and Society. Science, 2012, 336, 673-674.	12.6	39
86	Investigating the potential for ethnic group harm in collaborative genomics research in Africa: Is ethnic stigmatisation likely?. Social Science and Medicine, 2012, 75, 1400-1407.	3.8	38
87	Seeking consent to genetic and genomic research in a rural Ghanaian setting: A qualitative study of the MalariaGEN experience. BMC Medical Ethics, 2012, 13, 15.	2.4	97
88	"It's for a good cause, isn't it?" - Exploring views of South African TB research participants on sample storage and re-use. BMC Medical Ethics, 2012, 13, 19.	2.4	27
89	Editorial (An Idea Whose Time Has Come? An African Foresight Observatory on Genomics Medicine and) Tj ETQq1 1 0,784314 rgBT /Ove	0.2	11
90	Ethical, legal and social issues in the context of the planning stages of the Southern African Human Genome Programme. Medicine and Law, 2012, 31, 119-52.	0.0	12

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91	Assessing the Privacy Risks of Data Sharing in Genomics. <i>Public Health Genomics</i> , 2011, 14, 17-25.	1.0	99
92	Ethical issues in human genomics research in developing countries. <i>BMC Medical Ethics</i> , 2011, 12, 5.	2.4	120
93	Ethical implications of the use of whole genome methods in medical research. <i>European Journal of Human Genetics</i> , 2010, 18, 398-403.	2.8	98
94	Ethical Data Release in Genome-Wide Association Studies in Developing Countries. <i>PLoS Medicine</i> , 2009, 6, e1000143.	8.4	68
95	Public Access to Genome-Wide Data: Five Views on Balancing Research with Privacy and Protection. <i>PLoS Genetics</i> , 2009, 5, e1000665.	3.5	71
96	Data sharing in genomics “re-shaping scientific practice. <i>Nature Reviews Genetics</i> , 2009, 10, 331-335.	16.3	253
97	Planning for translational research in genomics. <i>Genome Medicine</i> , 2009, 1, 87.	8.2	6
98	The Path Forward for DNA Data. <i>Science</i> , 2008, 322, 1186-1187.	12.6	0
99	The obesity epidemic: medical and ethical considerations. <i>Science and Engineering Ethics</i> , 2007, 13, 55-67.	2.9	47
100	Using the Drama of DNA approach to community engagement in genomic research in South Africa: experiences and lessons learnt. <i>AAS Open Research</i> , 0, 3, 1.	1.5	5
101	Ethical challenges in community engagement practices in research during the COVID-19 pandemic in Africa. <i>AAS Open Research</i> , 0, 3, 23.	1.5	12
102	Policy makers, regulators and researchers’ perspectives on genomics research and the capacity of the National Health Research Act of 2013 to regulate genomics research in Zambia. <i>AAS Open Research</i> , 0, 3, 32.	1.5	3
103	Should institutions fund the feedback of individual findings in genomic research?. <i>Journal of Medical Ethics</i> , 0, , medethics-2021-107992.	1.8	2
104	Whole Exome Sequencing in South Africa: Stakeholder Views on Return of Individual Research Results and Incidental Findings. <i>Frontiers in Genetics</i> , 0, 13, .	2.3	5